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			Application Number	09/633,573	
INF	ORMATION	DISCLOSURE	Filing Date	08/04/2000	
STA	TEMENT 8	BY APPLICANT	First Named Inventor	Wilson T. Asfora	
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	(use as many s	heets as necessary)	Examiner Name		
Sheet	1	of 1	Attorney Docket Number	00-0050	

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Sheet	1	of	2

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Application Number	09/633,573		
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First Named Inventor	Wilson T. Asfora		
Group Art Unit			
Examiner Name			
Attorney Docket Number	00-0050		

				U.S. PATENT DOC	UMENTS	
Examiner Initials*	Cite No.1	U.S. Patent Number	Document Kind Code ² (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
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SM		5,683,357		Magram	11/04/1997	
∞		5,913,852		Magram	06/22/1999	
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	FOREIGN PATENT DOCUMENTS							
Examiner Cite			Foreign Patent Do	cument	Name of Patentee or	Date of Publication of	Pages, Columns, Lines, Where Relevant	
	No.1	Office ³	Number ⁴	Kind Code ⁵ (if known)	Applicant of Cited Document	Cited Document MM-DD-YYYY	Passages or Relevant Figures Appear	Ţθ

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Sheet 2 of 2

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First Named Inventor	Wilson T. Asfora		
Group Art Unit			
Examiner Name			
Attorney Docket Number	00.0050		

		OTHER PRIOR ART NON PATENT LITERATURE DOCUMENTS	
Examiner Initials*	Cite No.1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T²
m		Medical Printout of Premarket Notification (510(k)) for Medial Devices from U.S. Food and Drug Administration website for 510(k) number K974726	
Am		Hedical Printout of Premarket Notification (510(k)) for Medial Devices from U.S. Food and Drug Administration website for 510(k) number K970578	
SW		Printout of Premarket Notification (510(k)) for Medial Devices from U.S. Food and Drug Administration website for 510(k) number K984053	
SM		Medical Printout of Premarket Notification (510(k)) for Medial Devices from U.S. Food and Drug Administration website for 510(k) number K981046	
SM		Printout of Premarket Notification (510(k)) for Medial Devices from U.S. Food and Drug Administration website for 510(k) number K982702	
AN	•	Medical Printout of Premarket Notification (510(k)) for Medial Devices from U.S. Food and Drug Administration website for 510(k) number K962928	
m		Medical Printout of Premarket Notification (510(k)) for Medial Devices from U.S. Food and Drug Administration website for 510(k) number K981846	
BW		Copies from "Neurological and Neurosurgical Intensive Care", Third Edition. The article title is "Intracranial Pressure Monitoring Devices: Principles, Insertion, and Care". Consisting of pages 53-68. Dated 1993. Published by Raven Press, Ltd., 1185 Avenue of the Americans, New York, NY 10036.	
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